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FILE COVERS 1907 - 11 Sep 2008 VOL 149 ISS 11 FILE LAST UPDATED: 10 Sep 2008 (20080910/ED)

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Structure attributes must be viewed using STN Express query preparation. L2 727 SEA FILE=REGISTRY SSS FUL L1 L3

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ANSWER 1 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2008:831355 CAPLUS

DOCUMENT NUMBER: 149:152831

TITLE: Process for preparation of 2-amino-5-cyanobenzamides from the corresponding 5-halo compounds using metal cyanides in the presence of cuprous salts, iodides,

and amines.

Annis, Gary David; Bruening, Joerg; Currie, Martin INVENTOR(S): James; Dumas, Donald Joseph; Shapiro, Rafael

E. I. Du Pont de Nemours and Company, USA PATENT ASSIGNEE(S):

PCT Int. Appl., 63 pp. SOURCE:

CODEN: PIXXD2 DOCUMENT TYPE: Patent Enalish

LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PAT	ENT	NO.			KIN	D	DATE			APPL	ICAT	ION I	NO.		D	ATE	
WO	2008	0825	02		A2		2008	0710		WO 2	007-	US25	800		2	0071	218
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PRIORITY APPLN. INFO .:

US 2006-876394P US 2007-902465P

P 20061221 20070221

OTHER SOURCE(S):

CASREACT 149:152831; MARPAT 149:152831

GI

AB Title compds. (I; R1 = NHR3, OR4; R2 = Me, C1; R3 = H, alkyl, cyclopropyl, cyclopropylcyclopropyl, cyclopropylmethyl, methylcyclopropyl; R4 = H, alkyl), were prepared by treatment of the corresponding halides (II; Y = Br, C1; R1, R2 as above) with a metal cvanide in the presence of a Cu(I) salt, an iodide salt, and R5HNCR6R7(CR8R9)nCR10R11XR12 (X = NR13, O; n = 0, 1; R5, R7, R8, R9, R11, R12 = H, alkyl; R6, R10 = H, alkyl, Ph; R13 = H, Me; n = 0, 1). Thus, 2-amino-5-bromo-N, 3-dimethylbenzamide (preparation given), NaCN, CuI, and N, N'-dimethylethylenediamine were heated in xylenes at 140° for 4.5 h to give 2-amino-5-cyano-N,3-dimethylbenzamide.

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preparation); PREP (Preparation)
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RL: IMF (Industrial manufacture); PRPH (Prophetic); SPN (Synthetic
preparation); PREP (Preparation)
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(prepn of aminocyanobenzamides from the corresponding halo compds. using metal cyanides in the presence of cuprous salts, iodides, and amines)

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1042429-39-9P 1042429-40-2P
1042429-43-5P 1042429-44-6P
1042429-30-0P
               1042429-31-1P
1042429-37-7P
               1042429-38-8P
1042429-41-3P
               1042429-42-4P
                              1042429-43-5P 1042429-44-6P
1042429-50-4P 1042429-52-6P
1042429-47-9P 1042429-49-1P
RL: IMF (Industrial manufacture); PRPH (Prophetic); SPN (Synthetic
```

(prepn of aminocyanobenzamides from the corresponding halo compds.

(prepn of aminocyanobenzamides from the corresponding halo compds. using metal cyanides in the presence of cuprous salts, iodides, and amines)

L3 ANSWER 2 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:735755 CAPLUS

DOCUMENT NUMBER: 149:47043

TITLE: Synergistic insecticidal compositions comprising an

anthranilamide derivative

INVENTOR(S): Koyanagi, Toru; Morita, Masayuki; Yoneda, Tetsuo;
Ueda, Tsuvoshi; Kiriyama, Kazuhisa; Hamamoto, Taku

PATENT ASSIGNEE(S): Ishihara Sangyo Kaisha, Ltd., Japan

SOURCE: PCT Int. Appl., 82pp.

CODEN: PIXXD2
DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

GI

	PATENT NO.					D	DATE				ICAT				D	ATE	
						-									-		
WO	2008	0727:	83		A1		2008	0619		WO 2	007-	JP74:	372		2	0071	212
	W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BH,	BR,	BW,	BY,	BZ,	CA,
		CH,	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DO,	DZ,	EC,	EE,	EG,	ES,	FI,
		GB,	GD,	GE,	GH,	GM,	GT,	HN,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,
		KM,	KN,	KP,	KR,	KZ,	LA,	LC,	LK,	LR,	LS,	LT,	LU,	LY,	MA,	MD,	ME,
	MG, MK, M					MX,	MY,	MZ,	NA,	NG,	NI,	NO,	NZ,	OM,	PG,	PH,	PL,
	PT, RO, R					SC,	SD,	SE,	SG,	SK,	SL,	SM,	SV,	SY,	TJ,	TM,	TN,
	PT, RO, RI TR, TT, TI					UG,	US,	UZ,	VC,	VN,	ZA,	ZM,	ZW				
	RW:	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,
		IS,	IT,	LT,	LU,	LV,	MC,	MT,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,
		ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG,	BW,
		GH,	GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	AZ,
		BY,	KG,	KZ,	MD,	RU,	TJ,	TM									
PRIORITY	IORITY APPLN. INFO.:									JP 2	006-	3365	85		A 2	0061	214
										JP 2	007-	1050	29		A 2	0070	412
OTHER SO	HER SOURCE(S):						149:	4704	3								

AB Synergistic insecticidal and ectoparasiticidal compns. contain a anthranilamide derivs. I [Rla, Rlb = halo; R2, R3 = halo, alkyl, haloalkyl, alkoxy, haloalkoxy or cyano; A = alkyl substituted by Y; Y = (un)substituted C3-4 cycloalkyl; n = 0 or 1; q = 0, 1-4] and another pesticide.

Ι

REFERÈNCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT IT 1031756-99-6 1031757-01-3 1031757-03-5

T 1031756-99-6 1031757-01-3 1031757-03-5 1031757-05-7 1031757-07-9 1031757-10-4 1031757-14-8 1031757-17-1 1031757-19-3 1031757-22-1 7031757-23-9 1031757-26-2 1031757-28-4 1031757-30-8 1031757-32-0

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1031757-34-2 1031757-36-4 1031757-38-6
1031757-41-1 1031757-43-3 1031757-45-5
1031757-47-7 1031757-49-9 1031757-51-3
1031757-54-6 1031757-56-8 1031757-58-0
1031757-60-4 1031757-62-6 1031757-64-8
1031757-66-0 1031757-69-3 1031757-71-7
1031757-74-0 1031757-76-2 1031757-78-4
1031757-80-8 1031757-82-0 1031757-84-2
1031757-86-4 1031757-88-6 1031757-90-0
1031757-92-2 1031757-94-4
RL: AGR (Agricultural use); BUU (Biological use, unclassified); BIOL
(Biological study); USES (Uses)
   (synergistic insecticidal compn)
112410-23-8D, Tebufenozide, mixts. with anthranilamide derivs.
112839-32-4D, Furconazole-cis, mixts. with anthranilamide derivs.
114369-43-6D, Fenbuconazole, mixts. with anthranilamide derivs.
115852-48-7D, Fenoxanil, mixts. with anthranilamide derivs.
116255-48-2D, Bromuconazole, mixts. with anthranilamide derivs.
116714-46-6D, Novaluron, mixts. with anthranilamide derivs.
117428-22-5D, Picoxystrobin, mixts, with anthranilamide derivs.
118134-30-8D, Spiroxamine, mixts, with anthranilamide derivs.
119168-77-3D, Tebufenpyrad, mixts. with anthranilamide derivs.
119446-68-3D, DIfenoconazole, mixts. with anthranilamide derivs.
119544-94-4D, Protrifenbute, mixts. with anthranilamide derivs.
119791-41-2D, Emamectin, mixts. with anthranilamide derivs.
120068-37-3D, Fipronil, mixts. with anthranilamide derivs.
                                                            120116-88-3D,
Cvazofamid, mixts, with anthranilamide derivs, 120928-09-8D, Fenazaguin,
mixts. with anthranilamide derivs. 121451-02-3D, Noviflumuron, mixts.
with anthranilamide derivs. 121552-61-2D, Cyprodinil, mixts. with
anthranilamide derivs. 122453-73-0D, Chlorfenapyr, mixts. with
anthranilamide derivs. 123312-89-0D, Pymetrozine, mixts. with
anthranilamide derivs. 124495-18-7D, Quinoxyfen, mixts. with
anthranilamide derivs. 125116-23-6D, Metconazole, mixts. with
anthranilamide derivs. 125225-28-7D, Ipconazole, mixts. with
anthranilamide derivs. 126069-54-3D, Phosphocarb, mixts. with
anthranilamide derivs.
                       126833-17-8D, Fenhexamid, mixts. with
anthranilamide derivs.
                       129558-76-5D, Tolfenpyrad, mixts. with
anthranilamide derivs.
                       130000-40-7D, Thifluzamide, mixts. with
anthranilamide derivs.
                       131341-86-1D, Fludioxonil, mixts, with
anthranilamide derivs. 131807-57-3D, Famoxadone, mixts, with
anthranilamide derivs.
                       131860-33-8D, Azoxystrobin, mixts, with
anthranilamide derivs.
                       133855-98-8D, Epoxiconazole, mixts. with
anthranilamide derivs.
                        134098-61-6D, Fenpyroximate, mixts. with
anthranilamide derivs.
                        135410-20-7D, Acetamiprid, mixts. with
anthranilamide derivs.
                        136426-54-5D, Fluquinconazole, mixts, with
                        138261-41-3D, Imidacloprid, mixts, with
anthranilamide derivs.
                        139920-32-4D, Diclocymet, mixts, with
anthranilamide derivs.
anthranilamide derivs.
                        139968-49-3D, Metaflumizone, mixts, with
                        140163-89-9D, Imicyafos, mixts. with
anthranilamide derivs.
anthranilamide derivs.
                        140923-17-7D, Iprovalicarb, mixts. with
                        141517-21-7D, Trifloxystrobin, mixts. with
anthranilamide derivs.
                        143390-89-0D, Kresoximmethyl, mixts. with
anthranilamide derivs.
anthranilamide derivs.
                        143807-66-3D, Chromafenozide, mixts. with
anthranilamide derivs.
                        148477-71-8D, Spirodiclofen, mixts. with
anthranilamide derivs.
                       149508-90-7D, Sipconazole, mixts. with
anthranilamide derivs. 149877-41-8D, Bifenazate, mixts. with
anthranilamide derivs. 149961-52-4D, Dimoxystrobin, mixts. with
anthranilamide derivs. 150824-47-8D, Nitenpyram, mixts. with
anthranilamide derivs. 153233-91-1D, Etoxazole, mixts. with
anthranilamide derivs. 153719-23-4D, Thiamethoxam, mixts. with
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anthranilamide derivs.
                          155569-91-8D, Emamectin benzoate, mixts. with
anthranilamide derivs. 156052-68-5D, Zoxamide, mixts. with
anthranilamide derivs. 158062-67-0D, Flonicamid, mixts. with
anthranilamide derivs. 161050-58-4D, Methoxyfenozide, mixts. with
anthranilamide derivs. 161326-34-7D, Fenamidone, mixts. with
anthranilamide derivs. 162650-77-3D, Ethaboxam, mixts. with
anthranilamide derivs. 165252-70-0D, Dinotefuran, mixts. with
anthranilamide derivs. 168316-95-8D, Spinosad, mixts, with
anthranilamide derivs. 170015-32-4D, Flufenerim, mixts, with
anthranilamide derivs. 173584-44-6D, Indoxacarb, mixts. with
anthranilamide derivs. 174212-12-5D, Oxpoconazole fumarate, mixts. with
anthranilamide derivs.
                          175013-18-0D, Pyraclostrobin, mixts. with
anthranilamide derivs.
                           175217-20-6D, Silthiopham, mixts. with
anthranilamide derivs.
                          178928-70-6D, Prothioconazole, mixts. with
anthranilamide derivs. 179101-81-6D, Pyridalyl, mixts. with
anthranilamide derivs. 180409-60-3D, Cyflufenamid, mixts. with anthranilamide derivs. 181587-01-9D, Ethiprole, mixts. with anthranilamide derivs. 182916-02-5D, Metominofen, mixts. with anthranilamide derivs. 183675-82-3D, MTF-753, mixts. with anthranilamide
          188425-85-6D, Boscalid, mixts, with anthranilamide derivs.
189278-12-4D, Proquinazid, mixts, with anthranilamide derivs.
203313-25-1D, Spirotetramat, mixts. with anthranilamide derivs.
210880-92-5D, Clothianidin, mixts. with anthranilamide derivs.
211867-47-9D, Flumorph, mixts. with anthranilamide derivs. 220899-03-6D,
Metrafenone, mixts. with anthranilamide derivs. 223419-20-3D,
Profluthrin, mixts. with anthranilamide derivs.
                                                     223580-51-6D, Tiadinil,
mixts, with anthranilamide derivs. 229977-93-9D, Fluacrypyrim, mixts.
with anthranilamide derivs. 240494-70-6D, ,Metofluthrin, mixts. with
anthranilamide derivs. 248593-16-0D, Orysastrobin, mixts. with
anthranilamide derivs. 272451-65-7D, Flubendiamide, mixts. with
anthranilamide derivs. 283594-90-1D, Spiromesifen, mixts. with
anthranilamide derivs. 315208-17-4D, Pyrafluprole, mixts. with anthranilamide derivs. 337458-27-2D, Pyrifluquinazon, mixts. with
anthranilamide derivs. 348635-87-0D, Amisulbrom, mixts. with
anthranilamide derivs. 361377-29-9D, Fluoxastrobin, mixts. with
anthranilamide derivs. 394730-71-3D, Pyriprole, mixts. with anthranilamide derivs. 400882-07-7D, Cyflumetofen, mixts. with anthranilamide derivs. 413615-35-7D, Benthiavalicarb-, mixts. with
anthranilamide derivs. 500008-45-7D, Chlorantraniliprole, mixts. with
anthranilamide derivs. 560121-52-0D, Cyenopyrafen, mixts. with
anthranilamide derivs. 863549-51-3D, Lepimectin, mixts. with
anthranilamide derivs. 935545-74-7D, DE 175, mixts. with anthranilamide
derivs. 937279-54-4D, HGW 86, mixts. with anthranilamide derivs.
946494-18-4D, UBF 307, mixts. with anthranilamide derivs. 946494-19-5D,
KIF 7767, mixts. with anthranilamide derivs. 946494-20-8D, Syngenta
446510, mixts. with anthranilamide derivs. 1031756-98-5D, mixts.
containing 1031757-98-8D, mixts. containing 1031758-00-5D,
mixts. containing 1032110-39-6D, BCM 062, mixts. with anthranilamide derivs.
1032111-18-4D, BCM 061, mixts. with anthranilamide derivs.
1032111-42-4D, BCF 051, mixts. with anthranilamide derivs.
RL: AGR (Agricultural use); BUU (Biological use, unclassified); BIOL
(Biological study); USES (Uses)
    (synergistic insecticidal compns.)
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L3 ANSWER 3 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2008:734496 CAPLUS

DOCUMENT NUMBER: 149:79593

TITLE: Process for production of anthranilamide compound
INVENTOR(S): Koyanagi, Toru; Yamamoto, Kazuhiro; Yoneda, Tetsuo;
Kanbayashi, Shigehisa; Tanimura, Toyoshi; Taquchi,

Yohei; Yoshida, Tatsunori

PATENT ASSIGNEE(S): Ishihara Sangyo Kaisha, Ltd., Japan

SOURCE: PCT Int. Appl., 95pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: PATENT INFORMATION:

PA	PATENT NO.					D	DATE		1	APPL	ICAT	ION	NO.		D	ATE	
WO	2008				A1	-	2008	0619	1	WO 2	007-	JP74	169		2	0071	214
	W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BH,	BR,	BW,	BY,	BZ,	CA,
		CH,	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DO,	DZ,	EC,	EE,	EG,	ES,	FI,
		GB,	GD,	GE,	GH,	GM,	GT,	HN,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,
		KM,	KN,	KP,	KR,	KZ,	LA,	LC,	LK,	LR,	LS,	LT,	LU,	LY,	MA,	MD,	ME,
		MG,	MK,	MN,	MW,	MX,	MY,	MZ,	NA,	NG,	NI,	NO,	NZ,	OM,	PG,	PH,	PL,
		PT,	RO,	RS,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SM,	SV,	SY,	TJ,	TM,	TN,
		TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	ZA,	ZM,	ZW				
	RW:	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,
		IS,	IT,	LT,	LU,	LV,	MC,	MT,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,
		ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG,	BW,
		GH,	GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	AZ,
		BY,	KG,	KZ,	MD,	RU,	TJ,	TM									
PRIORIT	Y APP	LN.	INFO	. :						JP 2	006-	3391	00		A 2	0061	215
										JP 2	007-	1527	18		A 2	0070	808
OTHER S	DURCE	(S):			MAR	PAT	149:	7959	3								

AB Disclosed is a process for producing a specific anthranilamide compound or a salt thereof. Specifically disclosed is a process for producing an anthranilamide compound represented by the formula I: [wherein Rla and R3 independently represent a halogen or a haloalkyl; R2 represents a cyclopropylalkyl or a cyclobutylalkyl; and HaI represents a chlorine atom or a bromine atom] or a salt thereof, which comprises the step of selectively halogenating a compound represented by the formula I: [wherein Rla, R2 and R3 are as defined above, HaI = H]. For example, II was provided in a multi-step synthesis starting from the reaction of Et 2-furoylpyruvate with 3-chloro-2-hydrazylpyridine.

REFERENCE COUNT: 55 THERE ARE 55 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT IT 1621-24-5P, (1-Cyclopropylethyl)amine 51761-72-9P, Cyclopropylmethylketoxime 107855-32-3P 112881-69-3P 112881-76-2P

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1033344-64-7P, 1-(3-Chloropyridin-2-y1)-5-(2-fury1)-1H-pyrazole-3-
    carboxylic acid 1033344-84-1P, 3-[(Benzyloxycarbonyl)amino]-1-(3-
    chloropyridin-2-y1)-1H-5-pyrazolecarboxylic acid 1033344-87-4P,
    3-[(Benzyloxycarbonyl)amino]-1-(3-chloropyridin-2-yl)-5-(2-furyl)-1H-
    pyrazole 1033407-53-2P 1033407-55-4P 1033407-56-5P,
    N-[4-Chloro-2-(1-cyclopropylethylcarbamoyl)phenyl]-1-(3-chloropyridin-2-
    v1)-3-hvdroxv-4,5-dihvdro-1H-pvrazole-5-carboxamide 1033407-57-6P
    1033407-58-7P, 3-Bromo-N-[4-chloro-2-(1-cyclopropylethylcarbamoyl)phenyl]-
    1-(3-chloropyridin-2-yl)-4,5-dihydro-1H-pyrazole-5-carboxamide
    1033407-59-8P, 3-Bromo-N-[4-chloro-2-(1-
    cyclopropylethylcarbamoyl)phenyl]-1-(3-chloropyridin-2-yl)-1H-pyrazole-5-
    carboxamide 1033407-60-1P
                                 1033407-61-2P, 2-Amino-5-chloro-N-(1-
    cyclopropylethyl)benzamide
                                 1033407-63-4P 1033407-64-5P, Pentyl
    3-bromo-1-(3-chloropyridin-2-yl)-4,5-dihydro-1H-pyrazole-5-carboxylate
    1033407-65-6P, Pentyl 3-bromo-1-(3-chloropyridin-2-yl)-1H-pyrazole-5-
    carboxylate 1033407-66-7P, Phenyl 3-bromo-1-(3-chloropyridin-2-yl)-1H-
    pyrazole-5-carboxylate 1033407-67-8P, 2-Amino-3-bromo-5-chloro-N-(1-
    cyclopropylethyl)benzamide 1033407-69-0P, Ethyl 1-(3-chloropyridin-2-yl)-
    5-(2-furyl)-4,5-dihydro-1H-pyrazole-3-carboxylate 1033407-70-3P,
    3-[(Benzyloxycarbonyl)amino]-1-(3-chloropyridin-2-yl)-1H-5-
    pyrazolecarboxylic acid phenyl ester 1033407-72-5P, 3-Amino-1-(3-
    chloropyridin-2-yl)-1H-5-pyrazolecarboxylic acid phenyl ester
    1033407-73-6P, Benzyl 3-bromo-1-(3-chloropyridin-2-yl)-1H-pyrazole-5-
    carboxylate 1033407-75-8P 1033407-76-9P
                                                 1033407-78-1P.
    5-Chloro-N-(1-cyclopropylethyl)-2-nitrobenzamide
    RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
    (Reactant or reagent)
        (preparation of anthranilamide compound)
    1031756-98-5P, 3-Bromo-N-[2-bromo-4-chloro-6-[[(1-
    cyclopropylethyl)amino]carbonyl]phenyl]-1-(3-chloropyridin-2-yl)-1H-
                            1033407-74-7P, 4-Methoxybenzyl
    pyrazole-5-carboxamide
    3-bromo-1-(3-chloropyridin-2-yl)-1H-pyrazole-5-carboxylate
    RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation of anthranilamide compound)
    ANSWER 4 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                        2008:733566 CAPLUS
DOCUMENT NUMBER:
                        149:79590
TITLE:
                        Process for preparation of anthranilamide compound by
                        using novel pyrazole compound as intermediate
INVENTOR(S):
                        Koyanagi, Toru; Hisamatsu, Akihiro
PATENT ASSIGNEE(S):
                       Ishihara Sangyo Kaisha, Ltd., Japan
SOURCE:
                        PCT Int. Appl., 59pp.
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        Japanese
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:
    PATENT NO.
                        KIND
                               DATE
                                          APPLICATION NO.
                                                                 DATE
    WO 2008072743
                              20080619
                                        WO 2007-JP74166 20071214
                         A1
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA,
            CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI,
            GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG,
            KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME,
            MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL,
            PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN,
            TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
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RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,

IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

PRIORITY APPLN. INFO.:

JP 2006-339100 A 20061215 JP 2007-128991 A 20070515 JP 2007-137551 A 20070524

OTHER SOURCE(S):

MARPAT 149:79590

GI

AB Disclosed is a method for producing an anthranilamide compound or a salt thereof. Specifically disclosed is a method for producing an anthranilamide compound represented by the formula I [ R1 = halo, alkyl, alkenyl, etc.; A = (un)substituted alkyl; X = halo; m = 0-4] or a salt thereof, which is characterized in that a compound represented by the formula I [R1, A and m are defined as above; X = NH2] is diazotized and then reacted with copper halide, copper metal or an alkyl halide. For example, II was provided in a multi-step synthesis starting from the reaction of Et 2-furoylpyruvate with 3-chloro-2-hydrazylpyridine. REFERENCE COUNT: 48 THERE ARE 48 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

1031756-98-5P

RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)

(preparation of anthranilamide compound by using novel pyrazole compound as intermediate)

L3 ANSWER 5 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2008:703045 CAPLUS

DOCUMENT NUMBER: 149:53719

TITLE: Process for preparing 2-amino-5-cyanobenzoates by treatment of 2-amino-5-halobenzoates with alkali metal nitriles in the presence of palladium phosphine

catalysts.

INVENTOR(S): Bruening, Joerg; Casalnuovo, Albert Loren; Grushin,

Vladimir

PATENT ASSIGNEE(S): E. I. du Pont de Nemours and Company, USA

SOURCE: PCT Int. Appl., 50pp. CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

GI

## PATENT INFORMATION:

PATENT	NO.		KIN		DATE			APPL	ICAT					ATE	
WO 2008			A1			0612									
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PRIORITY APP	LN. INFO	. :						US 2	006-	8730	58P	1	P 2	0061	206
OTHER SOURCE	(S):		CASI	REAC'	T 14	9:53	719;	MAR	PAT :	149:	5371	9			

AB Title compds. (I, R1 = NHR3, OR4; R2 = Me, C1; R3 = H, alkyl, cyclopropyl, cyclopropylmethyl, methylcyclopropyl; R4 = H, alkyl; with a proviso); were prepared by treatment of (II; X = Br, C1; other variables as above) with MICN (M1 = alkali metal) in the presence of ≥1 ether and nitrile solvent, ≥1 palladium tertiary phosphine catalyst. Thus, 2-amino-5-bromo-N,3-dimethylbenzamide (preparation given), Zn, NaCN, and a catalyst solution prepared from tris (dibenzylideneacetone)dipalladium and tri-tert-butylphosphine were stirred together in THF for 63 h at 25° to give >99% conversion to 2-amino-5-cyano-N,3-

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dimethylbenzamide.
REFERENCE COUNT:
                               THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
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RL: IMF (Industrial manufacture); PRPH (Prophetic); SPN (Synthetic
preparation): PREP (Preparation)
     (preparation of aminocyanobenzoates by treatment of aminohalobenzoates with
     alkali metal nitriles in the presence of palladium phosphine catalysts)
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RL: IMF (Industrial manufacture); PRPH (Prophetic); SPN (Synthetic preparation); PREP (Preparation)

(preparation of aminocyanobenzoates by treatment of aminohalobenzoates with alkali metal nitriles in the presence of palladium phosphine catalysts)

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RL: IMF (Industrial manufacture); PRPH (Prophetic); SPN (Synthetic
preparation); PREP (Preparation)
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(preparation of aminocyanobenzoates by treatment of aminohalobenzoates with alkali metal nitriles in the presence of palladium phosphine catalysts)

ANSWER 6 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:673452 CAPLUS

DOCUMENT NUMBER: 149:10005

TITLE: Preparation of (heterocyclyl) N-

cyanoalkylanthranilamides as insecticides and

acaricides

Muehlebach, Michel; Craig, Gerald Wayne INVENTOR(S): PATENT ASSIGNEE(S): Syngenta Participations AG, Switz.

SOURCE: PCT Int. Appl., 91pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PRI

PAT	TENT :	NO.			KIN	D	DATE			APPL	ICAT	ION :	NO.		D	ATE	
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RITY	Y APP	LN.	INFO	. :						EP 2	006-	2486	5		A 2	0061	201
R SC	DURCE	(S):			MARI	PAT	149:	1000	5								

OTH GI

AB Title compds. [I; D = (substituted) Ph, pyridyl, pyrazolyl, pyrrolyl, pyridyl, pyrimidyl; n = 0-3; R1 = halo, OH, NO2, alkyl, alkenyl, alkynyl, cycloalkyl, alkylthio, cycloalkylamino, (substituted) Ph, PhCH2, PhO, etc.; R2, R3 = H, (substituted) alkyl, alkenyl, alkynyl, cycloalkyl; E1, E2 = O, S; R4 = (substituted) alkyl, cycloalkyl; R5 = (substituted) cycloalkyl, cycloalkylalkyl], were prepared Thus, title compound (II) was prepared in 4 steps from 2-amino-5-chloro-3-methylbenzoic acid, cyclopropyl Me ketone, and 2-(3-chloropyridin-2-vl)-5-trifluoromethyl-2H-pyrazole-3carbonyl chloride. II and other I at 200 ppm gave >80% control of Spodoptera littoralis. REFERENCE COUNT: THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT TT 1034824-68-4P 1034824-71-9P 1034824-74-2P 1034824-77-5P 1034824-80-0P 1034824-82-2P 1034824-84-4P 1034824-87-7P 1034824-90-2P 1034824-93-5P 1034824-96-8P 1034824-99-1P 1034825-02-9P 1034825-05-2P 1034825-08-5P 1034825-11-0P 1034825-14-3P 1034825-17-6P 1034825-20-1P 1034825-23-4P 1034825-26-7P 1034825-29-0P 1034825-32-5P 1034825-35-8P 1034825-38-1P 1034825-41-6P 1034825-44-9P 1034825-47-2P 1034825-50-7P 1034825-53-0P 1034825-56-3P 1034825-59-6P 1034825-62-1P 1034825-65-4P 1034825-68-7P 1034825-72-3P 1034825-75-6P 1034825-78-9P 1034825-81-4P 1034825-84-7P 1034825-87-0P 1034825-90-5P 1034825-93-8P 1034825-96-1P 1034825-99-4P 1034826-02-2P 1034826-05-5P 1034826-08-8P 1034826-11-3P 1034826-14-6P 1034826-17-9P 1034826-20-4P 1034826-23-7P 1034826-26-0P 1034826-29-3P 1034826-32-8P 1034826-35-1P 1034826-38-4P 1034826-41-9P 1034826-44-2P 1034826-47-5P 1034826-50-0P 1034826-52-2P 1034826-54-4P 1034826-56-6P 1034826-59-9P 1034826-62-4P 1034826-64-6P 1034826-67-9P 1034826-70-4P 1034826-73-7P 1034826-75-9P 1034826-78-2P 1034826-81-7P 1034826-84-0P 1034826-86-2P 1034826-89-5P 1034826-92-0P 1034826-95-3P 1034826-97-5P 1034826-99-7P 1034827-01-4P 1034827-04-7P 1034827-07-0P 1034827-09-2P 1034827-11-6P 1034827-13-8P 1034827-15-0P 1034827-17-2P 1034827-19-4P 1034827-23-0P 1034827-25-2P 1034827-27-4P 1034827-29-6P 1034827-31-0P 1034827-33-2P 1034827-36-5P 1034827-38-7P 1034827-41-2P 1034827-44-5P 1034827-47-8P 1034827-49-0P 1034827-51-4P 1034827-53-6P 1034827-55-8P 1034827-58-1P 1034827-60-5P 1034827-62-7P 1034827-64-9P 1034827-66-1P 1034827-69-4P 1034827-71-8P 1034827-73-0P 1034827-75-2P 1034827-77-4P 1034827-79-6P 1034827-81-0P 1034827-83-2P 1034827-85-4P 1034827-87-6P 1034827-89-8P 1034827-91-2P 1034827-93-4P 1034827-95-6P 1034827-97-8P 1034827-99-0P 1034828-95-6P 1034827-97-8P 1034828-05-1P 1034828-07-3P 1034828-05-1P 1034828-07-3P 1034828-07-3P 1034828-07-3P 1034828-13-1P 1034828-13-1P 1034828-15-3P 1034828-15-3P 1034828-19-1P 1034828-11-1P 1034828-19-3P 1034828-27-3P 1034828-27-3P 1034828-27-3P 1034828-37-9P 1034828-31-3P 1034828-37-3P 1034828-37-9P 1034828-37-3P 1034828-38-3P 1034828-38-3P 1034828-38-3P 1034828-38-3P 1034828-38-3P 1034828-3P 1034828-

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RL: AGR (Agricultural use); BSU (Biological study, unclassified); PRPH
(Prophetic); SPN (Synthetic preparation); BIOL (Biological study); PREP
(Preparation); USES (Uses)
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RL: AGR (Agricultural use); BSU (Biological study, unclassified); PRPH
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(Prophetic); SPN (Synthetic preparation); BIOL (Biological study); PREP
     (Preparation); USES (Uses)
        (preparation of (heterocyclyl) N-cyanoalkylanthranilamides as insecticides
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     (Preparation); USES (Uses)
        (preparation of (heterocyclyl) N-cyanoalkylanthranilamides as insecticides
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        (preparation of (heterocyclyl) N-cyanoalkylanthranilamides as insecticides
        and acaricides)
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     1029975-12-9P 1029975-14-1P 1029975-16-3P
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     RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN
    (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (preparation of (heterocyclyl) N-cyanoalkylanthranilamides as insecticides
        and acaricides)
L3 ANSWER 7 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                        2008:221501 CAPLUS
DOCUMENT NUMBER:
                         148:231873
TITLE:
                        Crop vigor and yield enhancement and
                        arthropod-vectored plant disease disruption by
                        carboxamide derivatives
INVENTOR(S):
                        Annan, Isaac Billy; Marcon, Paula Cristina Rodrigues
                        Gouveia; Portillo, Hector Eduardo
PATENT ASSIGNEE(S):
                        E. I. Du Pont de Nemours and Company, USA
SOURCE:
                        PCT Int. Appl., 23pp.
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
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	WO	2008	0211	52		A2	_	2008	0221		WO 2	007-	US17	673			0070	
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			CH,	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DO,	DZ,	EC,	EE,	EG,	ES,	FΙ,
			GB,	GD,	GE,	GH,	GM,	GT,	HN,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,
			KM,	KN,	KP,	KR,	ΚZ,	LA,	LC,	LK,	LR,	LS,	LT,	LU,	LY,	MA,	MD,	ME,
	MG, MK, MN, PT. RO. RS.			MN,	MW,	MX,	MY,	MZ,	NA,	NG,	NI,	NO,	NZ,	OM,	PG,	PH,	PL,	
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PRIO	PRIORITY APPLN. INFO.:										US 2	006-	8368	92P	1	P 2	0060	809
											US 2	006-	8370.	59P	1	P 2	0060	810
											US 2	006-	8504	54P	1	P 2	0061	010
	HED COUDCE (C).																	

OTHER SOURCE(S): MARPAT 148:231873 GI

- AB The anthranilamide anthropodicides I (X = N, CF, CCl, CBr or CI; Rl = Me, Cl, Br or F; R2 = H, F, Cl, Br or CN; R3 = F, Cl, Br, Cl-4 haloalkyl or haloalkow; R4a = H, Cl-4 alkyl cyclopropylmethyl or 1-cyclopropylethyl; R4b = H or Me; R5, r6 = H, F, Cl or Br) or their N-oxides, as well as phthalic acid derivs. (Markush given) enhance crop vigor and crop yield and disrupt infectious disease transmission by arthropod pests.
- IT 272451-65-7 438450-41-0 500008-00-4 500008-44-6 500008-45-7 736994-60-8 736994-63-1 736995-23-6 871238-02-7 871238-03-8 871238-04-9 882401-50-5 886583-54-6

886583-69-3

RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses) (crop vigor and yield enhancement and arthropod-vectored plant disease disruption by)

L3 ANSWER 8 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:91157 CAPLUS

DOCUMENT NUMBER: 148:191926

TITLE: Process for making 3-substituted 2-amino-5-

halobenzamides

INVENTOR(S): Davis, Richard Frank; Shapiro, Rafael; Taylor, Eric

Deguyon

PATENT ASSIGNEE(S): E. I. du Pont de Nemours and Company, USA

SOURCE: PCT Int. Appl., 38pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE:

English FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA:	ENT:	ΝΟ.			KIN	D	DATE			APPL	ICAT	ION	NO.		D	ATE	
WO	2008				A2	_	2008	0124		WO 2	007-	US14	972		2	0070	627
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		CH,	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DO,	DZ,	EC,	EE,	EG,	ES,	FI,
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RITY	APP	LN.	INFO	. :						US 2	006-	8317	81P	1	P 2	0060	719
R SC	URCE	(S):			CASI	REAC	T 14	8:19	1926	; MAI	RPAT	148	:191	926			

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## \* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

AB Disclosed is a method for preparing I (R1 = H, alkyl, cyclopropyl, cyclopropylmethyl, or methylcyclopropyl; R2 = Me or C1; X = C1 or Br) by ring opening of II with R1-NH2 in the presence of a carboxylic acid and a method for preparing II by cyclization of III (R3 = (un)substituted alkyl or alkenyl) with phosphorus tribromide. Also disclosed is a method for preparing known insecticides IV (R4 = C1, Br, CF3, OCF2H or OCH2CF3; Z = CR7

		wii Ilisectitides i			
		Cl or $Br$ ; $R6 = F$		H, F, Cl or B	
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 RL: PRPH (Prophetic); SPN (Synthetic preparation); PREP (Preparation)
        (preparation of aminoarylcarboxamides via PBr3 induced cyclization of
        carboxamidobenzoic acids followed by ring opening with alkyl amines)
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RL: PRPH (Prophetic); SPN (Synthetic preparation); PREP (Preparation)
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(preparation of aminoarylcarboxamides via PBr3 induced cyclization of carboxamidobenzoic acids followed by ring opening with alkyl amines)

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L3 ANSWER 9 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
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ACCESSION NUMBER: 2007:1278752 CAPLUS

DOCUMENT NUMBER: 147:481495

TITLE: Use of carboxamide derivatives for disrupting the

reproductive performance of arthropods INVENTOR(S): Annan, Isaac Billy; Flexner, John Lindsey; Marcon,

Paula Cristina Rodrigues Gouveia; Portillo, Hector

Eduardo

E. I. Du Pont De Nemours and Company, USA PATENT ASSIGNEE(S):

SOURCE: PCT Int. Appl., 34pp. CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

PATENT NO.	KIND DA	DATE APPLI	ICATION NO.	DATE
WO 2007126636	A2 20	20071108 WO 20	007-US6929	20070320
WO 2007126636	A3 20	20080320		
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WO 2007-US6929 W 20070320
PRIORITY APPLN. INFO.:
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OTHER SOURCE(S): MARPAT 147:481495

GΙ

AB The carboxamide derivs. I (X = n, CF, CCl, CBr or CI; R1 = Me, Cl, Br or F; R2 = H, F, C1, Br or CN; R3 = F, C1, Br, haloalkyl or haloalkoxy; R4a = H, alkyl, cyclopropylmethyl or 1-cyclopropylethyl; R4b = H or Me; R5, R6 = H,F, Cl or Br), their N-oxides and salts.

272451-65-7 438450-41-0 500008-00-4 500008-44-6 500008-45-7 736994-60-8 736994-63-1 736995-23-6 871238-02-7 871238-03-8 871238-04-9 882401-50-5 886583-54-6

Ι

886583-69-3

RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses) (use of carboxamide derivs. for disrupting the reproductive performance of arthropods)

ANSWER 10 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2006:1120573 CAPLUS

DOCUMENT NUMBER: 145:455006

TITLE: Preparation of cyanoanthranilamides as insecticides

and acaricides

Jeanquenat, Andre; O'Sullivan, Anthony; Muehlebach, INVENTOR(S):

Michel; Trah, Stephan; Hall, Roger Graham PATENT ASSIGNEE(S): Syngenta Participations AG, Switz.

SOURCE: PCT Int. Appl., 100pp. CODEN: PIXXD2

DOCUMENT TYPE: Pat.ent.

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

> PATENT NO. KIND DATE APPLICATION NO. DATE

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WO 2006111341
                        A1 20061026 WO 2006-EP3504
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PRIORITY APPLN. INFO.:
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OTHER SOURCE(S):
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AB Title compds. [I, E, Z = 0, S; A = (substituted) alkylene, alkenylene, alkynylene, bivalent mono- or bicyclic ring; X = 0, NH, alkylimino; Y = (substituted) mono- or bicyclic ring; p, q = 0, I; B = (substituted) 3-4 membered (heterocyclic) ring; R1 = halo, NO2, cyano, OH, alkyl, alkenyl, alkynyl, cycloalkyl, haloalkyl, (substituted) Ph, PhCH2, PhO, etc., n = 0-3; R2, R3 = H, alkyl, alkenyl, alkynyl, substituted cycloalkyl; D = (substituted) Ph, pyridyl, pyrradyl, pyrmidyl), yprimylyl, vere prepared Thus, 2-[2-(3-chloropyridin-2-yl)-5-trifluoromethyl-2H-pyrazol-3-yl]-8-methyl-4-oxo-4H-benzo[d] [1,3]oxazine-6-carbonitrile, bicycloprop-1-ylamine hydrochloride (preparation given), and Et3N were heated together in THF at 60° for 8 h to give 2-(3-chloropyridin-2-yl)-5-trifluoromethyl-2H-pyrazole-3-carboxylic acid [2-(bicycloprop-1-ylcarbamoyl)]-4-cyano-6-methylphenyl]amide. The latter at 400 ppm showed >80% activity against Cydia pomonella.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT IT 1042422-55-8 1042425-11-5 1042425-51-3 1042426-12-9 1042426-58-3

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RL: PRPH (Prophetic)
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(Preparation of cyanoanthranilamides as insecticides and acaricides)

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L3 ANSWER 11 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                     2006:558556 CAPLUS
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DOCUMENT NUMBER:

145:62886 TITLE:

Anthranilamide derivatives as insecticides, and their preparation, pesticidal compositions and formulation INVENTOR(S): Jeanguenat, Andre; O'Sullivan, Anthony Cornelius
PATENT ASSIGNEE(S): Syngenta Participations A.-G., Switz.
SOURCE: PCT Int. Appl., 136 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

LANGUAGE: Er FAMILY ACC. NUM. COUNT: 1

Patent English

FAMILY ACC. NUM. COUNT PATENT INFORMATION:

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			SG,	SK,	SL,	SM,	SY,	TJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,
			VN,	YU,	ZA,	ZM,	ZW											
		RW:	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,
			IS,	IT,	LT,	LU,	LV,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,	BJ,
			CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG,	BW,	GH,
			GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	ΑZ,	BY,
			KG,	KZ,	MD,	RU,	TJ,	TM										
		2005																
	EP	1819	695			A1		2007	0822		EP 2	2005-	8154	27		2	0051	207
		R:										ES,						
			IS,	IT,	LI,	LT,	LU,	LV,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR	
		1010																
	JP	2008	5230	8 0		T		2008	0703		JP 2	2007-	5448	09		2	0051	207
	US	2008	0146	552		A1		2008	0619		US 2	2007-	7205	71		2	0070	531
	IN	2007	DN 04	178		A		2007	0831		IN 2	2007-	DN41	78		2	0070	601
	MX	2007	0689	8		A		2007	0626		MX 2	2007-	6898			2	0070	608
		2007				A		2007	0904									
PRIO	IORITY APPLN. INFO.:											2004-						
												2005-				W 2	0051	207
	R S	DURCE	(S):			CAS	REAC	T 14	5:62	886;	MAF	RPAT	145:	6288	6			
GI																		

\* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT \*

Compds. of formula I, and the agrochem. acceptable salts and all stereoisomers and tautomeric forms of the compds. of formula I can be used as agrochem, active ingredients and can be prepared in a manner known per se. Several examples on formulation of compds. of formula I is also disclosed in this invention. Compds. of formula I wherein El and W2 are independently O or S; R1 is halo, CN, NO2, OH, C1-6 (halo)alkyl, C2-6 (halo)alkenyl, C2-6 (halo)alkynyl, C3-6 (halo)cycloalkyl, C1-4 (halo) alkoxy, C1-4 (halo) alkylthio, C1-4 (halo) alkylsulfinyl, C1-4 (halo)alkylsulfonyl, C1-4 alkylamino, C2-4 dialkylamino, C3-6 cycloalkylamino, etc.; n is 0, 1, 2, 3, or 4; R2 and R3 are independently H, (un)substituted C1-6 alkvl, (un)substituted C2-6 alkenvl, (un) substituted C2-6 alkynyl, or (un) substituted C3-6 cycloalkyl; D is (un) substituted Ph, (un) substituted pyridyl, (un) substituted pyrazole, (un) substituted pyrrole, or (un) substituted pyrimidine; Yla and Y2 are independently (un) substituted C1-6 alkylene, (un) substituted C2-6 alkenylene, or (un)substituted C3-6 alkynylene, etc.; G is a bond, O, N-Z1, S or G1-C(=G2)-G3; G1 and G3 are independently a bond, O, S, or NZ2; G2 is O, S or NZ3; Z and Z1-Z3 are independently H, C1-6 (halo)alkyl, C2-6 (halo)alkenyl, C2-6 (halo)alkynyl, C3-6 (halo)cycloalkyl, C1-4 (halo)alkoxy, C1-4 (halo)alkylthio, etc.; Y3 is H, halo or C1-6

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(halo)alkyl; Ylb is a bond, or (un)substituted C1-6 alkylene,
     (un) substituted C2-6 alkenylene, or (un) substituted C3-6 alkynylene; and
     their tautomers, agrochem. utilizable salts and auxiliary are claimed.
     Example compound II was prepared by amidation of 6-chloro-2-[2-(3-
     chloropyridin-2-yl)-5-trifluoromethyl-2H-pyrazol-3-yl]-8-
     methylbenzo[d][1,3]oxazin-4-one with 1-amino-2-propanol; the resulting
     2-(3-chloropyridin-2-vl)-5-trifluoromethyl-2H-pyrazol-3-carboxylic acid
     [4-chloro-2-(2-hydroxypropylcarbamovl)-6-methylphenyl]amide underwent
     substitution with thioacetic acid to give thioacetic acid
     S-[2-(5-chloro-2-[[2-(3-chloropyridin-2-yl)-5-trifluoromethyl-2H-pyrazole-
     3-carbonyl]amino]-3-methylbenzoylamino)-1-methylethyl] ester, which
     underwent deacetylation and methylation to give the corresponding Me thio
     ether, which underwent oxidation to give the corresponding sulfoxide, which
     reacted with trifluoroacetamide to give the corresponding
     N-trifluoroacetylated sulfoximide, which underwent deacetylation to give
     compound II. All the invention compds. were evaluated for their
     insecticidal activity. Some of the tested compds. showed good activity
     against Aphis craccivora, Diabrotica balteata, Heliothis virescens
     (application to foliar and egg), Myzus persicae (foliar and systemic
     application), Plutella xvlostella and Spodoptera littoralis.
REFERENCE COUNT:
                               THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
                                  891195-06-5P 891195-07-6P 891195-08-7P
891195-11-2P 891195-12-3P 891195-13-4P
     891195-04-3P 891195-05-4P
     891195-09-8P 891195-10-1P
     891195-14-5P 891195-15-6P 891195-16-7P 891195-17-8P
891195-18-9P 891195-19-0P 891195-20-3P 891195-21-4P
     891195-22-5P 891195-23-6P 891195-25-8P 891195-26-9P 891195-27-0P 891195-28-1P 891195-29-2P 891195-30-5P 891195-31-6P 891195-32-7P
     891195-33-8P 891195-34-9P 891195-35-0P 891195-36-1P 891195-37-2P
     891195-38-3P
     RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN
     (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES
     (Uses)
        (agrochem. candidate; preparation of anthranilamide derivs. as insecticides)
   ANSWER 12 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                        2006:496102 CAPLUS
DOCUMENT NUMBER:
                         144:462625
TITLE:
                         Preparation of anthranilamide derivative insecticides
                         and acaricides
INVENTOR(S):
                         Lahm, George Philip; Selby, Thomas Paul; Stevenson,
                         Thomas Martin; Taggi, Andrew Edmund; Bereznak, James
                         Francis
PATENT ASSIGNEE(S):
                        E.I. Dupont De Nemours and Co., USA
                        PCT Int. Appl., 97 pp.
SOURCE:
                         CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
     PATENT NO.
                        KIND DATE APPLICATION NO. DATE
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                                                                    _____
                     A2
                        A2 20060526
A3 20061221
     WO 2006055922
                                          WO 2005-US42196
     WO 2006055922
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
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CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX,
MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE,
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SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC,
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             CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH,
             GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
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                                20060526
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     AU 2005306363
                          A1
                                                                   20051118
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                          A1
                                20060526
                                            CA 2005-2585378
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     EP 1812421
                          A2
                                20070801
                                            EP 2005-851952
                                                                   20051118
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             BA, HR, MK, YU
     CN 101061103
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     IN 2007DN03224
                          Α
                                20070831
                                            IN 2007-DN3224
                                                                   20070430
                                            KR 2007-713584
     KR 2007086280
                          Α
                                20070827
                                                                   20070615
PRIORITY APPLN. INFO .:
                                            US 2004-629120P
                                                                P 20041118
                                                               P 20050610
                                            US 2005-689414P
                                                               W 20051118
                                            WO 2005-US42196
                       MARPAT 144:462625
OTHER SOURCE(S):
GΙ
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NR2-COJ CONR3R4 I

AB The anthranilamide derivs. I and their geometric and stereoisomers,

N-oxides, and salts [J = (un)substituted Ph or N-containing heterocyclyl; Rl =
alkyl alkenyl, alkynyl, etc.; R2 = alkylcarbonyl, alkoxycarbonyl or
(di)alkylaminocarbonyl; R3 = (cyclo)alkyl, alkenyl, alkynyl, alkoxy, etc.
; R4 = (un)substituted alkylcycloalkyl, alkenyl, alkynyl, cycloalkyl,
alkynylcycloalkyl, cycloalkylalkyl, cycloalkylalkenyl, cycloalkylalkynyl,
cycloalkenylalkyl or alkylcycloalkenyl, oxiranylalkyl, thiiranylalkyl,
oxetanylalkyl, thietanylalkyl, 3-oxetanyl or 3-thietanyl; R5 =
(cyclo)alkyl, haloalkyl, alkenyl alkynyl, etc.] are prepared as pesticides
for controlling invertebrate pests, specifically insecticides and
acaricides.

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736995-23-6P 882401-50-5P 886583-28-4P
886583-29-5P 886583-30-8P 886583-31-9P
                                            886583-32-0P
                                                          886583-33-1P
886583-34-2P
              886583-35-3P
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886583-39-7P
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886583-54-6P 886583-55-7P
                            886583-56-8P
                                            886583-57-9P
886583-58-0P 886583-59-1P 886583-69-3P
RL: AGR (Agricultural use); SPN (Synthetic preparation); BIOL (Biological
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study); PREP (Preparation); USES (Uses)

(preparation as insecticide and acaricides)

IT 886583-65-9 886583-66-0 886583-67-1

886583-68-2

RL: AGR (Agricultural use); BIOL (Biological study); USES (Uses)

(synergistic insecticide and acaricide)

L3 ANSWER 13 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2006:367128 CAPLUS

DOCUMENT NUMBER: 144:364548

TITLE: Preparation of anthranilamide derivative acaricides

and insecticides

INVENTOR(S): O'Sullivan, Anthony Cornelius; Hughes, Dave;

Jeanguenat, Andre; Muehlebach, Michel; Loiseleur,

Olivier

PATENT ASSIGNEE(S): Syngenta Participations AG, Switz.; Syngenta Limited SOURCE: PCT Int. Appl., 152 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

	TENT											ICAT:					ATE	
	2006																	
	2006																	
	W:	AE.	AG.	AL.	AM.	AT.	AU.	AZ,	BA.	BE	3.	BG.	BR.	BW.	BY.	BZ.	CA.	CH.
								DK,										
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	RW:					CY.	CZ.	DE,	DK.	EB	٥.	ES.	FI.	FR.	GB.	GR.	HU.	IE.
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								SD,										
					RU.			,	,		•	,		,		,	,	,
AU	2005	2938	01		A1		2006	0420		AU	20	005-2	29380	01		2	0051	010
	2580																	
	1802																	
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CN	1010	6111	0		A		2007	1024		CN	20	005-1	30039	9682		2	0051	010
JP	2008	5158	44		T		2008	0515		JΡ	20	007-	5351:	14		2	0051	010
IN	2007	DN02	167		A		2007	0803		IN	20	007-1	DN21	67		2	0070	320
MX	2007 2007	0390	8		A		2007	0521		MX	20	007-3	3908			2	0070	330
KR	2007	0635	36		A		2007	0619		KR	20	007-	7082	49		2	0070	411
RIORIT	Y APP	LN.	INFO	. :						GB	20	004-2	22556	6		A 2	0041	011
										WO	20	005-E	EP108	891	1	W 2	0051	010
PHED C	OTTROE	101.			C7 C1	0070	T 14	1.36	15.10	. 1.	47\ T	TEGG	144	. 261	5.10			

OTHER SOURCE(S): CASREACT 144:364548; MARPAT 144:364548

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$$\begin{array}{c|c} E & D \\ \hline & N & R^2 \\ \hline & Z \\ \hline & N-AX_pY_qB \\ \hline & R^3 & I \end{array}$$

The anthranilamides I [E, Z = 0 or S; A, Y = alkylene, alkenylene,ΔR alkynylene, etc.; X = O, NH or alkyl-substituted NH; B = (un)substituted ring; D = (un)substituted Ph, pyridyl, pyrazolyl, etc.; R1 = amino, formyl, cyanoalkenyl, etc.; R2, R3 = H, (un)substituted alkyl, alkenyl, cycloalkyl, etc.; n = 0, 1-4; p, q = 0 or 1] and I salts, stereoisomers and tautomers are prepared as acaricides and insecticides. 882401-41-4P 882401-42-5P 882401-43-6P 882401-44-7P 882401-45-8P 882401-46-9P 882401-47-0P 882401-48-1P 882401-49-2P 882401-50-5P 882401-51-6P 882401-52-7P 882401-53-8P 882401-54-9P 882401-55-0P 882401-56-1P 882401-57-2P 882401-58-3P 882401-59-4P 882401-60-7P 882401-61-8P 882401-62-9P 882401-63-0P 882401-64-1P 882401-65-2P 882401-66-3P 882401-67-4P 882401-68-5P 882401-69-6P 882401-70-9P 882401-71-0P 882401-72-1P 882401-73-2P 882401-74-3P 882401-75-4P 882401-76-5P 882401-77-6P 882401-78-7P 882401-79-8P 882401-80-1P 882401-81-2P 882401-82-3P 882401-83-4P 882401-84-5P 882401-85-6P 882401-86-7P 882401-87-8P 882401-88-9P 882401-89-0P 882401-90-3P 882401-91-4P 882401-92-5P 882401-93-6P 882401-94-7P 882401-95-8P 882401-96-9P 882401-97-0P 882401-98-1P 882401-99-2P 882402-00-8P 882402-01-9P 882402-02-0P 882402-03-1P 882402-04-2P 882402-05-3P 882402-06-4P 882402-07-5P 882402-08-6P 882402-09-7P 882402-10-0P RL: AGR (Agricultural use); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation as acaricide and insecticide) ANSWER 14 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:902883 CAPLUS

DOCUMENT NUMBER: 143:229846

TITLE: Preparation of anthranilamides as pesticides

INVENTOR(S): Koyanagi, Toru; Morita, Masayuki; Nakamoto, Kenichi;

Hisamatsu, Akihiro

PATENT ASSIGNEE(S): Ishihara Sangvo Kaisha, Ltd., Japan

PCT Int. Appl., 52 pp. SOURCE: CODEN: PIXXD2

DOCUMENT TYPE: Pat.ent. LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

> PATENT NO. KIND DATE APPLICATION NO. DATE

GI

WO	2005	0779	34		A1		2005	0825		WO	2005-	JP23	51		2	0050	216
	W:	AE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB	, BG,	BR,	BW,	BY,	BZ,	CA,	CH,
		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ	, EC,	EE,	EG,	ES,	FI,	GB,	GD,
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS	, JP,	KE,	KG,	KP,	KR,	KZ,	LC,
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		NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU	, SC,	SD,	SE,	SG,	SK,	SL,	SY,
		TJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US	, UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW
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		ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	ТJ,	TM,	AT	, BE,	BG,	CH,	CY,	CZ,	DE,	DK,
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	2005										2005-						
	2553				A1						2005-					0050	216
EP	1717										2005-					0050	
	R:										, IT,						
						FΙ,	RO,	MK,	CY,	ΑL	, TR,	BG,	CZ,	EE,	HU,	PL,	SK,
			HR,	IS,													
	1918				A		2007				2005-					0050	
	2005						2007				2005-					0050	
	2006						2007				2006-					0060	
	2006						2006				2006-					0060	
	2007				A1		2007	0607			2006-					0060	
PRIORITY	Y APP	LN.	INFO	. :							2004-					0040	
											2004-					0040	
											2004-					0040	
											2004-					0041	
										WO	2005-	JP23	51		W 2	0050	216
OTHER SO	JURCE	(S):			MAR	PAT	143:	2298	16								

$$(\mathbb{R}^1)_m \xrightarrow{\begin{subarray}{c} H & \square \\ N-C & N \\ 0 & (\mathbb{O})_n \end{subarray}} \mathbb{R}^2$$

AB The title anthranilamides, i.e. N-(2-aminocarbonylphenyl)-1-(2-puyla(J)-1H-pyrazole-5-carboxamide derivs. represented by the general formula (I) or
salts thereof [wherein Rl = halogeno, alkyl, haloalkyl, alkenyl,
haloalkenyl, alkynyl, haloalkynyl, alakoxy, haloalkoxy, alkylcarbonyl,
haloalkylcarbonyl, NO2, CHO; R2, R3 = halogeno, alkyl, haloalkyl, alkoxy,
haloalkoxy, cyano; A = Y-substituted alkyl (Y = C3-4 cycloalkyl optionally
substituted by ≥1 groups selected from halogeno, alkyl, and
haloalkyl); n = 0,1; q = 0-4; provided that R1 is F, C1, Br, or Me
substituted at 2-position of the benzene ring and another R1 is halogeno
substituted at 4-position of the benzene ring the 4-halogeno group is F
or C1] are prepared They are useful as pesticides, in particular
insecticides, acaricides, nematocides, and parasticides. Thus, 1.49 g

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Et3N was slowly added dropwise to a solution of 0.8 g cyclopropylmethylamine
    hydrochloride in 40 mL THF, stirred at room temperature for 30 min, slowly
    treated dropwise with a solution of 1 g 2-[1-(3-chloro-2-pyridy1)-3-
    (trifluoromethyl)-1H-pyrazol-5-yl]-8-methyl-4H-3,1-benzoxazin-4-one in 10
    mL THF, and refluxed for 4 h to give, after workup and silica gel
    chromatog., 0.54 g N-[6-[[(cyclopropylmethyl)amino]carbonyl]-2-
    methylphenyl]-1-(3-chloro-2-pyridyl)-3-(trifluoromethyl)-1H-pyrazole-5-
    carboxamide (II). II at 3.1 ppm controlled 2-nd to 3-rd instar larvae of
    Spodoptera litura on cabbage leaves.
REFERENCE COUNT:
                        62
                              THERE ARE 62 CITED REFERENCES AVAILABLE FOR THIS
                              RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
    862995-50-4P 862995-51-5P 862995-52-6P
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    862995-86-6P 862995-87-7P 862995-88-8P
    RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN
    (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES
    (Uses)
       (preparation of anthranilamides as pesticides such as insecticides,
       acaricides, nematocides, and parasiticides)
    ANSWER 15 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER: 2004:648522 CAPLUS
DOCUMENT NUMBER:
                        141:190786
TITLE:
                       Preparation of cyano anthranilamide insecticides
INVENTOR(S):
                       Hughes, Kenneth Andrew; Lahm, George Philip; Selby,
                        Thomas Paul; Stevenson, Thomas Martin
PATENT ASSIGNEE(S):
                       E.I. Du Pont De Nemours and Company, USA
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DOCUMENT TYPE:
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LANGUAGE:
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FAMILY ACC. NUM. COUNT: 1
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                        A1 20040812 WO 2004-US3568 20040121
    WO 2004067528
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
            AU 2004207848
    CA 2512242
    EP 1599463
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
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PRIORITY APPLN. INFO.:			US	2003-443256P	P	20030128
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OTHER SOURCE(S):	MARPAT	141:190786				

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The title compds. [I; R1 = Me, C1, Br, F; R2 = F, C1, Br, haloalkyl or haloalkoxy; R3 = F, C1, Br; R4 = H, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, each optionally substituted with one substituent selected from the group consisting of halo, CN, SMe S(O)Me, S(O)2Me and OMe; R5 = H, Me; R6 = H, F, C1; R7 = H, F, C1], useful for controlling an invertebrate pest, were prepared E.g., a multi-step synthesis of compound I  $[R1=Me;\ R2=CF3;\ R3=C1;\ R4,\ R5=H],\ was given.$  The compds. I were tested in various biol. tests (data given). This invention also pertains to a composition for controlling an invertebrate pest comprising a biol. effective amount of a compound I, an N-oxide thereof or a suitable salt of the compound I and at least one addnl. component selected from the group consisting of a surfactant, a solid diluent and a liquid diluent.

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RL: AGR (Agricultural use); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of cyano anthranilamide insecticides)

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